



ATTORNEY DOCKET NUMBER: 2002834-0058 (CIP4 DIV/Modified Allergens)

**IN THE UNITED STATES PATENT AND TRADEMARK OFFICE**

Applicant: Bannon, et al.  
Serial No.: 09/478,668  
Filed: January 6, 2000  
For: METHODS AND REAGENTS FOR DECREASING CLINICAL REACTIONS  
TO ALLERGY

Examiner: Huynh, P.  
Art Unit: 1644

Assistant Commissioner for Patents  
Washington, DC 20231  
BOX AF

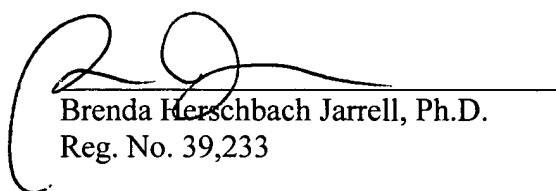
**TRANSMITTAL**

Enclosed please find the following documents regarding the above-referenced matter:

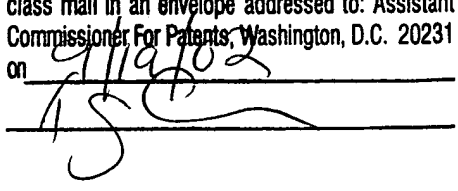
1. Supplemental Response to Office Action (3 page);
2. Exhibits A-G; and
3. a Return Postcard.

Please charge any fees or credit any overpayments to our Deposit Account No. 03-1721.

Respectfully submitted,

  
Brenda Herschbach Jarrell, Ph.D.  
Reg. No. 39,233

Choate, Hall & Stewart  
Exchange Place  
53 State Street  
Boston, MA 02109  
(617) 248-5000  
Dated: September 17, 2002  
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Applicant:	Bannon, et al.	Examiner:	Huynh, P.
Serial No.:	09/478,668	Art Unit:	1644
Filed:	January 6, 2000	Group:	1640
For:	METHODS AND REAGENTS FOR DECREASING CLINICAL REACTIONS TO ALLERGY		

#19  
JW  
10/2/02

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Washington, DC 20231  
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**SUPPLEMENTAL RESPONSE TO OFFICE ACTION**

Sir:

In response to the Final Office Action mailed December 18, 2001, the Interview held June 19, 2002 and further to the Response mailed June 18, 2002, Applicant respectfully requests consideration of the following Remarks.

**Remarks**

Applicant's representatives, Drs. Brenda Jarrell and Charles Lyon, would like to take this opportunity to thank Examiners Huynh and Chan for taking the time to hold an in person Interview on June 19, 2002. During the Interview, the Examiners indicated that they would be willing to consider evidence in support of Applicant's enablement argument. In particular, they indicated that they would consider post-art references that showed that the methods taught in the present application have been successfully applied to other allergens.

Accordingly, Applicant has collected a series of references showing that, after the present invention was made, people of ordinary skill in the art followed the steps taught in the present application (i.e., used patient sera to identify IgE binding epitopes, modified the protein sequence to alter identified IgE binding epitopes; and screened modified proteins to identify those with reduced binding) and were able without undue experimentation to obtain a variety of modified